IMPERIAL CANCER RESEARCH FUND LABORATORIES

M.G.P. STOKER, M.D., F.R.S. DIRECTOR OF RESEARCH

P.O. BOX NO. 123 LINCOLN'S INN FIELDS, LONDON, WC2A 3PX

Cables: Cancerch Tel.: 01-242 0200

Dr. Paul Berg,
Department of Biochemistry,
Stanford University Medical Center,
Stanford,
California 94305,
U.S.A.

1st February, 1973

Dear Paul,

This is just a brief progress report. You should by now have had the large batch of ST6 cells for testing. I hope they arrived safely. I also asked Bill to arrange for you to get a fresh batch of BHK clone C13 and to send you literature on the spiral bottles cell production system.

The transplantation antigen on ST1 cells is at the moment under test again. This time I have included an SV40 transformed BHK as a control as well as ST6.

Dr. Andy Louie, the postdoc who seemed to me to be a good person to repeat the isolation of abortive clones and to bring them to you for testing, has now opted for a different problem, namely a study on the histone-like proteins in the polyoma virus capsid and their relationship to the DNA. He is moving to Lionel to do this. I did not feel like dissuading him since the abortives project may not be so interesting now after all. However, during the last month or so he has isolated about ten abortive and stable transformants from a methocel transformation. They are in the freezer in case we want to do them later.

Now about the supposed partial genomes in ST1 and ST2 (and ST6?). All the ST series came from a single plate in one experiment. The

cells were layers of BHK which had been held in low ser/um and which were then infected by the method Joyce Taylor was using for the interferon studies. After four days in low serum they were suspended and plated for colonies. On one plate with 4×10^5 cells there were 75 transformed foci on a background of normal cells. 6 of these (ST1-6) were picked, grown and stored.

It might be something to do with the low serum, or it might be keeping the infected cells at high density for several days before replating and thus allowing opportunity for re-infections from the lytic cycles. Anyway, I have now arranged for this to be repeated as soon as possible. We will isolate 5 more clones in the ST series, grow them into large bat ches and if you are willing and interested will send them to you for testing, as well as to Vittorio Defendi for T-antigen.

I hope that there is some progress with Helene Smith's repeat experiment.

Geoff Clarke is now back after the Biohazard Meeting which he found valuable. I am very grateful to you for letting him come, and it will be good to see the record of the proceedings.

The reports of the ICRF departments for the review will be ready in a few weeks and we will be sending them on to you by air mail around the end of this month. It will be good to see you in May. The Queen is visiting the laboratories at about that time so perhaps she should join the review committee.

All the best,

Yours,

I starting from scratch but wifeling low come alls.